IN THE UNITED STATES DISTRICT COURT FOR THE WESTERN DISTRICT OF PENNSYLVANIA

IN RE: PHILIPS RECALLED CPAP,

Master Docket: Misc. No. 21-mc-1230-JFC

BI-LEVEL PAP, AND MECHANICAL

MDL No. 3014

LITIGATION

. 14152110.00

This Document Relates to:

VENTILATOR PRODUCTS

SHORT FORM COMPLAINT FOR PERSONAL INJURIES, DAMAGES,

HEATHER S. KEZDY, Individually and as Independent Administrator of the Estate of

AND DEMAND FOR JURY TRIAL

PIERRE G. KEZDY, deceased.

Plaintiff(s) incorporate(s) by reference the Amended Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial filed in *In re Philips Recalled CPAP, Bi-Level PAP, and Mechanical Ventilator Products Litigation*, MDL No. 3014, Master Docket Misc. No. 21-mc-1230 (the "Master Long Form Complaint"). This Short Form Complaint adopts the allegations, claims, and requested relief as set forth in the Master Long Form Complaint. As necessary herein, Plaintiff(s) may include: (a) additional claims and allegations against Defendants; and/or (b) additional claims and allegations against other Defendants not listed in the Master Long Form Complaint.

Plaintiff(s) further allege(s) as follows:

I. DEFENDANTS

- 1. Plaintiff(s) name(s) the following Defendants in this action:
 - Koninklijke Philips N.V.
 - Philips North America LLC.
 - Philips RS North America LLC.

II.

		Philips Holding USA Inc.
		Philips RS North America Holding Corporation.
		Polymer Technologies, Inc.
		Polymer Molded Products LLC.
II.	PLAI	NTIFF(S)
	2.	Name of Plaintiff(s):
		HEATHER S. KEZDY, Individually and as Independent Administrator of the Estate of PIERRE G. KEZDY, deceased
	3.	Name of spouse of Plaintiff (if loss of consortium claim is being made): HEATHER S. KEZDY
		HEATHER S. REZDT
	4.	Name and capacity (i.e., executor, administrator, guardian, conservator, etc.) of other Plaintiff, if any:
		Administrator
	5.	State(s) of residence of Plaintiff(s) (if the Recalled Device user is deceased, residence at the time of death): Illinois
III.	DESI	GNATED FORUM
	6.	Identify the forum (United States District Court and Division) in which the Plaintiff would have filed in the absence of direct filing:
		IN THE CIRCUIT COURT OF COOK COUNTY, ILLINOIS COUNTY DEPARTMENT, LAW DIVISION

IV. USE OF A RECALLED DEVICE

7. Plaintiff used the following Recalled Device(s):

□ E30 (Emergency Use Authorization) □ Dorma 500 □ DreamStation ASV □ REMstar SE Auto □ DreamStation ST, AVAPS □ Trilogy 100 □ SystemOne ASV4 □ Garbin Plus, Aeris, LifeVent □ C-Series ASV □ Garbin Plus, Aeris, LifeVent □ C-Series S/T and AVAPS □ A-Series BiPAP Hybrid A30 (not marketed in U.S.) □ SystemOne (Q-Series) □ A-Series BiPAP V30 Auto □ DreamStation □ A-Series BiPAP A40 □ DreamStation Go □ A-Series BiPAP A30 □ Dorma 400 □ Other Philips Respironics Device; if other, identify the model: □ REMstar CPAP machine (REF: 460P SN: P155811265313) with the attached System One HT Humid, DOM heated humidifier (REF: DS6TFLG V. INJURIES □ Remain (REF: 460P SN: P155811265313) with the attached System One HT Humid, DOM heated humidifier (REF: DS6TFLG V. INJURIES □ COPD (new or worsening) □ Asthma (new or worsening) □ Pulmonary Fibrosis □ Other Pulmonary Damage/Inflammatory Response □ Cancer □ (specify cancer)
DreamStation ST, AVAPS SystemOne ASV4 C-Series ASV Garbin Plus, Aeris, LifeVent A-Series BiPAP Hybrid A30 (not marketed in U.S.) SystemOne (Q-Series) A-Series BiPAP V30 Auto DreamStation DreamStation DreamStation Go Dorma 400 Dorma 400 New York Philips Respironics Device; if other, identify the model: REMstar CPAP machine (REF: 460P SN: P155811265313) with the attached System One HT Humid, DOM heated humidifier (REF: DS6TFLG) V. INJURIES 8. Plaintiff alleges the following physical injuries as a result of using a Recall-Device together with the attendant symptoms and consequences associated therewith: COPD (new or worsening) Asthma (new or worsening) Pulmonary Fibrosis ✓ Other Pulmonary Damage/Inflammatory Response
SystemOne ASV4 □ C-Series ASV □ Garbin Plus, Aeris, LifeVent □ C-Series S/T and AVAPS □ A-Series BiPAP Hybrid A30 (not marketed in U.S.) □ SystemOne (Q-Series) □ DreamStation □ DreamStation Go □ Dorma 400 □ DreamStation Go □ Other Philips Respironics Device; if other, identify the model: REMstar CPAP machine (REF: 460P SN: P155811265313) with the attached System One HT Humid, DOM heated humidifier (REF: DS6TFLG V. INJURIES 8. Plaintiff alleges the following physical injuries as a result of using a Recall Device together with the attendant symptoms and consequences associated therewith: □ COPD (new or worsening) □ Asthma (new or worsening) □ Pulmonary Fibrosis □ Other Pulmonary Damage/Inflammatory Response
□ C-Series ASV □ Garbin Plus, Aeris, LifeVent □ C-Series S/T and AVAPS □ A-Series BiPAP Hybrid A30 (not marketed in U.S.) □ SystemOne (Q-Series) □ A-Series BiPAP V30 Auto □ DreamStation □ A-Series BiPAP A40 □ Dorma 400 □ Other Philips Respironics Device; if other, identify the model: REMstar CPAP machine (REF: 460P SN: P155811265313) with the attached System One HT Humid, DOM heated humidifier (REF: DS6TFLG V. INJURIES 8. Plaintiff alleges the following physical injuries as a result of using a Recall-Device together with the attendant symptoms and consequences associated therewith: □ COPD (new or worsening) □ Asthma (new or worsening) □ Pulmonary Fibrosis ☑ Other Pulmonary Damage/Inflammatory Response
□ C-Series S/T and AVAPS □ A-Series BiPAP Hybrid A30 (not marketed □ OmniLab Advanced + in U.S.) □ SystemOne (Q-Series) □ A-Series BiPAP V30 Auto □ DreamStation □ A-Series BiPAP A40 □ Dorma 400 □ Other Philips Respironics Device; if other, identify the model: REMstar CPAP machine (REF: 460P SN: P155811265313) with the attached System One HT Humid, DOM heated humidifier (REF: DS6TFLG V. INJURIES 8. Plaintiff alleges the following physical injuries as a result of using a Recall-Device together with the attendant symptoms and consequences associated therewith: □ COPD (new or worsening) □ Asthma (new or worsening) □ Pulmonary Fibrosis ☑ Other Pulmonary Damage/Inflammatory Response
OmniLab Advanced +
SystemOne (Q-Series) A-Series BiPAP V30 Auto DreamStation A-Series BiPAP A40 DreamStation Go A-Series BiPAP A30 Other Philips Respironics Device; if other, identify the model: REMStar CPAP machine (REF: 460P SN: P155811265313) with the attached System One HT Humid, DOM heated humidifier (REF: DS6TFLG V. INJURIES 8. Plaintiff alleges the following physical injuries as a result of using a Recall Device together with the attendant symptoms and consequences associated therewith: COPD (new or worsening) Asthma (new or worsening) Pulmonary Fibrosis Other Pulmonary Damage/Inflammatory Response
DreamStation DreamStation Go A-Series BiPAP A30 Dorma 400 REMstar CPAP machine (REF: 460P SN: P15581 1265313) with the attached System One HT Humid, DOM heated humidifier (REF: DS6TFLG) V. INJURIES 8. Plaintiff alleges the following physical injuries as a result of using a Recall Device together with the attendant symptoms and consequences associated therewith: COPD (new or worsening) Asthma (new or worsening) Pulmonary Fibrosis ✓ Other Pulmonary Damage/Inflammatory Response
DreamStation Go Dorma 400 A-Series BiPAP A30 Dother Philips Respironics Device; if other, identify the model: REMstar CPAP machine (REF: 460P SN: P155811265313) with the attached System One HT Humid, DOM heated humidifier (REF: D86TFLG) V. INJURIES 8. Plaintiff alleges the following physical injuries as a result of using a Recall Device together with the attendant symptoms and consequences associated therewith: COPD (new or worsening) Asthma (new or worsening) Pulmonary Fibrosis ✓ Other Pulmonary Damage/Inflammatory Response
Dorma 400 ☐ Other Philips Respironics Device; if other, identify the model: REMstar CPAP machine (REF: 460P SN: P155811265313) with the attached System One HT Humid, DOM heated humidifier (REF: DS6TFLG) V. INJURIES 8. Plaintiff alleges the following physical injuries as a result of using a Recall Device together with the attendant symptoms and consequences associated therewith: ☐ COPD (new or worsening) ☐ Asthma (new or worsening) ☐ Pulmonary Fibrosis ✔ Other Pulmonary Damage/Inflammatory Response
identify the model: REMstar CPAP machine (REF: 460P SN: P155811265313) with the attached System One HT Humid, DOM heated humidifier (REF: DS6TFLG) V. INJURIES 8. Plaintiff alleges the following physical injuries as a result of using a Recall Device together with the attendant symptoms and consequences associated therewith: COPD (new or worsening) Asthma (new or worsening) Pulmonary Fibrosis Other Pulmonary Damage/Inflammatory Response
REMstar CPAP machine (REF: 460P SN: P155811265313) with the attached System One HT Humid, DOM heated humidifier (REF: DS6TFLG) V. INJURIES 8. Plaintiff alleges the following physical injuries as a result of using a Recall-Device together with the attendant symptoms and consequences associated therewith: COPD (new or worsening) Asthma (new or worsening) Pulmonary Fibrosis Other Pulmonary Damage/Inflammatory Response
P155811265313) with the attached System One HT Humid, DOM heated humidifier (REF: DS6TFLG) V. INJURIES 8. Plaintiff alleges the following physical injuries as a result of using a Recall Device together with the attendant symptoms and consequences associated therewith: COPD (new or worsening) Asthma (new or worsening) Pulmonary Fibrosis Other Pulmonary Damage/Inflammatory Response
8. Plaintiff alleges the following physical injuries as a result of using a Recall Device together with the attendant symptoms and consequences associated therewith: COPD (new or worsening) Asthma (new or worsening) Pulmonary Fibrosis Other Pulmonary Damage/Inflammatory Response
Kidney Damage
Liver Damage

VI.

	Heart Damage	
	✓ Death	
	Other (specify)	
CAUS	SES OF ACTION/DA	MAGES
9.	in the Master Long F	ilips N.V., Plaintiff(s) adopt(s) the following claims asserted form Complaint for Personal Injuries, Damages and Demand e allegations and prayer for relief with regard thereto, as set
	Count I:	Negligence
	Count II:	Strict Liability: Design Defect
	Count III:	Negligent Design
	Count IV:	Strict Liability: Failure to Warn
	Count V:	Negligent Failure to Warn
	Count VI:	Negligent Recall
	Count VII:	Battery
	Count VIII:	Strict Liability: Manufacturing Defect
	Count IX:	Negligent Manufacturing
	Count X:	Breach of Express Warranty
	Count XI:	Breach of the Implied Warranty of Merchantability
	Count XII:	Breach of the Implied Warranty of Usability
	Count XIII:	Fraud
	Count XIV:	Negligent Misrepresentation

Count XV:	Negligence Per Se
Count XVI:	Consumer Fraud and/or Unfair and Deceptive Practices Under State Law
Count XVII:	Unjust Enrichment
Count XVIII:	Loss of Consortium
Count XIX:	Survivorship and Wrongful Death
Count XX:	Medical Monitoring
Count XXI:	Punitive Damages
Count XXII:	Other [specify below]
asserted in the Maste	America LLC, Plaintiff(s) adopt(s) the following claims or Long Form Complaint for Personal Injuries, Damages and l, and the allegations and prayer for relief with regard thereto
Count I:	Negligence
Count II:	Strict Liability: Design Defect
Count III:	Negligent Design

Strict Liability: Failure to Warn

Strict Liability: Manufacturing Defect

Negligent Failure to Warn

Negligent Recall

Battery

10.

Count IV:

✓ Count V:

Count VI:

Count VII:

Count VIII:

Count IX:

Negligent Manufacturing

Count X:	Breach of Express Warranty
Count XI:	Breach of the Implied Warranty of Merchantability
Count XII:	Breach of the Implied Warranty of Usability
Count XIII:	Fraud
Count XIV:	Negligent Misrepresentation
Count XV:	Negligence Per Se
Count XVI:	Consumer Fraud and/or Unfair and Deceptive Practices Under State Law
Count XVII:	Unjust Enrichment
Count XVIII:	Loss of Consortium
Count XIX:	Survivorship and Wrongful Death
Count XX:	Medical Monitoring
Count XXI:	Punitive Damages
Count XXII:	Other [specify below]

11. As to Philips RS North America LLC, Plaintiff(s) adopt(s) the following claims asserted in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial, and the allegations and prayer for relief with regard thereto, as set forth therein:

Count I: Negligence

Count II: Strict Liability: Design Defect

Count III: Negligent Design

Count IV: Strict Liability: Failure to Warn

Count V:	Negligent Failure to Warn
Count VI:	Negligent Recall
Count VII:	Battery
Count VIII:	Strict Liability: Manufacturing Defect
Count IX:	Negligent Manufacturing
Count X:	Breach of Express Warranty
Count XI:	Breach of the Implied Warranty of Merchantability
Count XII:	Breach of the Implied Warranty of Usability
Count XIII:	Fraud
Count XIV:	Negligent Misrepresentation
Count XV:	Negligence Per Se
Count XVI:	Consumer Fraud and/or Unfair and Deceptive Practices Under State Law
Count XVII:	Unjust Enrichment
Count XVIII:	Loss of Consortium
Count XIX:	Survivorship and Wrongful Death
Count XX:	Medical Monitoring
Count XXI:	Punitive Damages
Count XXII:	Other [specify below]

12.	in the Master Long I	g USA Inc., Plaintiff(s) adopt(s) the following claims asserted form Complaint for Personal Injuries, Damages and Demand he allegations and prayer for relief with regard thereto, as set
	Count I:	Negligence
	Count II:	Strict Liability: Design Defect
	Count III:	Negligent Design
	Count IV:	Strict Liability: Failure to Warn
	Count V:	Negligent Failure to Warn
	Count VI:	Negligent Recall
	Count VII:	Battery
	Count VIII:	Strict Liability: Manufacturing Defect
	Count IX:	Negligent Manufacturing
	Count X:	Breach of Express Warranty
	Count XI:	Breach of the Implied Warranty of Merchantability
	Count XII:	Breach of the Implied Warranty of Usability
	Count XIII:	Fraud
	Count XIV:	Negligent Misrepresentation
	Count XV:	Negligence Per Se
	Count XVI:	Consumer Fraud and/or Unfair and Deceptive Practices Under State Law
	Count XVII:	Unjust Enrichment
	Count XVIII:	Loss of Consortium
	Count XIX:	Survivorship and Wrongful Death
	Count XX:	Medical Monitoring

Punitive Damages

Count XXI:

Count XIV:

Count XV:

	Count XXII:	Other [specify below]
13.	following claims asso	orth America Holding Corporation, Plaintiff(s) adopt(s) the erted in the Master Long Form Complaint for Personal Injuries, and for Jury Trial, and the allegations and prayer for relief with forth therein:
	Count I:	Negligence
	Count II:	Strict Liability: Design Defect
	Count III:	Negligent Design
	Count IV:	Strict Liability: Failure to Warn
	Count V:	Negligent Failure to Warn
	Count VI:	Negligent Recall
	Count VII:	Battery
	Count VIII:	Strict Liability: Manufacturing Defect
	Count IX:	Negligent Manufacturing
	Count X:	Breach of Express Warranty
	Count XI:	Breach of the Implied Warranty of Merchantability
	Count XII:	Breach of the Implied Warranty of Usability
	Count XIII:	Fraud

Negligence Per Se

Negligent Misrepresentation

Count XVI:	Consumer Fraud and/or Unfair and Deceptive Practices Under State Law
Count XVII:	Unjust Enrichment
Count XVIII:	Loss of Consortium
Count XIX:	Survivorship and Wrongful Death
Count XX:	Medical Monitoring
Count XXI:	Punitive Damages
Count XXII:	Other [specify below]

14. As to Polymer Technologies, Inc., Plaintiff(s) adopt(s) the following claims asserted in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial, and the allegations and prayer for relief with regard thereto, as set forth therein:

as set forth therein:	
Count I:	Negligence
Count II:	Strict Liability: Design Defect
Count III:	Negligent Design
Count IV:	Strict Liability: Failure to Warn
Count V:	Negligent Failure to Warn
Count VIII:	Strict Liability: Manufacturing Defect
Count IX:	Negligent Manufacturing
Count XIII:	Fraud
Count XIV:	Negligent Misrepresentation
Count XVII:	Unjust Enrichment

Count XVIII:	Loss of Consortium
Count XIX:	Survivorship and Wrongful Death
Count XX:	Medical Monitoring
Count XXI:	Punitive Damages
Count XXII:	Other [specify below]
Demand for Jury Trias set forth therein: Count I:	ter Long Form Complaint for Personal Injuries, Damages and ial, and the allegations and prayer for relief with regard thereto, Negligence
Count II:	Strict Liability: Design Defect
Count III:	Negligent Design
Count IV:	Strict Liability: Failure to Warn
Count V:	Negligent Failure to Warn
Count VIII:	Strict Liability: Manufacturing Defect
Count IX:	Negligent Manufacturing
Count XIII:	Fraud
Count XIV:	Negligent Misrepresentation

15.

Count XVII:

Count XVIII:

Count XIX:

Count XX:

Unjust Enrichment

Loss of Consortium

Medical Monitoring

Survivorship and Wrongful Death

Count XXI:	Punitive Damages
Count XXII:	Other [specify below]
Complaint for Personabove, the addition Plaintiff(s) assert(s	against the Defendants identified in the Master Long Form onal Injuries, Damages and Demand for Jury Trial are alleged al facts, if any, supporting these allegations must be pleaded.) the following additional factual allegations against the ed in the Master Long Form Complaint for Personal Injuries, and for Jury Trial:

18. Plaintiff(s) assert(s) the following additional claims and factual allegations against other Defendants named in Paragraph 16 above:

WHEREFORE, Plaintiff(s) pray(s) for relief and judgment against Defendants and all such further relief that this Court deems equitable and just as set forth in the Master Long Form Complaint for Personal Injuries, Damages and Demand for Jury Trial and any additional relief to which Plaintiff(s) may be entitled.

 /s/: Bryan Jambois

Bryan S. Jambois (bjambois@kjs-law.com) Steve K. Jambois (sjambois@kjs-law.com) KRALOVEC, JAMBOIS & SCHWARTZ 60 W. Randolph Street, 4th Floor Chicago, IL 60601 (312) 782-2525 Firm ID: 24797